Docket No.: PF-0484 US

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Printed:

February 1999



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of:

Hillman et al.

Title:

KINESIN LIGHT CHAIN HOMOLOG

Serial No.:

09/036,614

Filing Date:

March 6, 1998

Examiner:

Gucker, S.

Group Art Unit:

1645

Assistant Commissioner for Patents Washington, D.C. 20231

RESPONSE TO NOTICE TO COMPLY

Sir:

In response to the Examiner's communication and Notice to Comply mailed December 2, 1998, and in accordance with the requirements of 37 CFR § 1.821-1.825, Applicants hereby submit a substitute diskette containing the computer-readable information for the Sequence Listing of the above-identified application. The diskette complies with the requirements of 37 CFR § 1.824 and is IBM PC compatible using FastSEQ for Windows Version 2.0.

Accompanying the substitute diskette containing the computer-readable information is a substitute paper copy of the Sequence Listing as disclosed in the application.

The content of the substitute Sequence Listing paper copy is identical to the substitute computerreadable copy, as required under 37 CFR § 1.821(f).

Applicants believe no fee is due. If the Patent Office determines that a fee is due the Commissioner is authorized to debit Incyte Pharmaceuticals, Inc. Deposit Account No. 09-0108.

This response is enclosed in duplicate.

Respectfully submitted,

INCYTE PHARMACEUTICALS, INC.

Date: February 19

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37003

09/036,614

Application No.:09/036,614
NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING
MOSE EOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825 for the following reason(s):

X	 This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to these regulations, published at 1114 OG 29, May 15, 1990 and at 55 FR 18230, May 1, 1990.
	2. This application does not contain, as a separate part of the disclosure on paper copy, a "Sequence Listing" as required by 37 C.F.R. 1.821(c).
	3. A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e).
	4. A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up "Raw Sequence Listing."
X	5. The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d).
	6. The paper copy of the "Sequence Listing" is not the same as the computer readable from of the "Sequence Listing" as required by 37 C.F.R. 1.821(e).
	7. Other: Applicant Must Provide: An initial or <u>substitute</u> computer readable form (CRF) copy of the "Sequence Listing".
X	An initial or <u>substitute</u> paper copy of the "Sequence Listing", as well as an amendment directing its entry into the specification.
X	A statement that the content of the paper and computer readable copies are the same and, where applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).
For	questions regarding compliance to these requirements, please contact:
For	Rules Interpretation, call (703) 308-4216 CRF Submission Help, call (703) 308-4212 PatentIn software help, call (703) 308-6856

PLEASE RETURN A COPY OF THIS NOTICE WITH YOUR RESPONSE